

over the intended duration and environment of use;

(ii) Labeling must include the clinical training, if needed, for the safe use of this device and information on the patient population for which the device has been demonstrated to be effective;

(iii) For devices that incorporate electrical components, appropriate analysis and testing must validate electrical safety and electromagnetic compatibility;

(iv) For devices containing software, software verification, validation, and hazard analysis must be performed;

(v) Any elements of the device that may contact the patient device must be demonstrated to be biocompatible; and

(vi) For over-the-counter devices, human factors testing and analysis must validate that the device design and labeling are sufficient for lay use.

(c) *Premarket notification.* The CPR aid device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter if it is a prescription use device that provides feedback to the rescuer consistent with the current American Heart Association guidelines for CPR and in compliance with the special controls under paragraph (b)(2) of this section, subject to the limitations of exemptions in § 870.9.

Dated: January 2, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

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ARCHITECTURAL AND TRANSPORTATION BARRIERS COMPLIANCE BOARD

36 CFR Part 1195

[Docket No. ATCB-2012-0003]

RIN 3014-AA40

Medical Diagnostic Equipment Accessibility Standards Advisory Committee

AGENCY: Architectural and Transportation Barriers Compliance Board.

ACTION: Notice of advisory committee meeting.

SUMMARY: The Medical Diagnostic Equipment Accessibility Standards Advisory Committee will hold its third meeting. On July 5, 2012, the Architectural and Transportation Barriers Compliance Board (Access Board) established the advisory committee to make recommendations to

the Board on matters associated with comments received and responses to questions included in a previously published Notice of Proposed Rulemaking (NPRM) on Medical Diagnostic Equipment Accessibility Standards.

DATES: The Committee will meet on January 22, 2013, from 10:00 a.m. to 5:00 p.m. and on January 23, 2012, from 9:00 a.m. to 2:30 p.m.

ADDRESSES: The meeting will be held at the Access Board's Conference Room, 1331 F Street NW., Suite 800, Washington, DC 20004-1111.

FOR FURTHER INFORMATION CONTACT: Rex Pace, Office of Technical and Information Services, Architectural and Transportation Barriers Compliance Board, 1331 F Street NW., Suite 1000, Washington, DC 20004-1111. Telephone number (202) 272-0023 (Voice); (202) 272-0052 (TTY). Electronic mail address: pace@access-board.gov.

SUPPLEMENTARY INFORMATION: On July 5, 2012, the Architectural and Transportation Barriers Compliance Board (Access Board) established an advisory committee to make recommendations to the Board on matters associated with comments received and responses to questions included in a previously published NPRM on Medical Diagnostic Equipment Accessibility Standards. See 77 FR 6916 (February 9, 2012). The NPRM and information related to the proposed standards are available on the Access Board's Web site at: <http://www.access-board.gov/medical-equipment.htm>.

The advisory committee will hold its third meeting on January 22 and 23, 2013. The agenda includes the following:

- Review of previous committee work;
- Presentations by medical practitioners and clinicians on the use of medical diagnostic equipment in relation to transfer surfaces;
- Continued discussion on subcommittees based on medical diagnostic equipment type;
- Continued discussion on transfer surface height and size;
- Review and discussion on transfer support location and configuration;
- Consideration of issues proposed by committee members; and
- Discussion of administrative issues.

The preliminary meeting agenda, along with information about the committee, is available at the Access Board's Web site (<http://www.access-board.gov/medical-equipment.htm>).

Committee meetings are open to the public and interested persons can attend the meetings and communicate their views. Members of the public will have opportunities to address the committee on issues of interest to them during public comment periods scheduled on each day of the meeting.

The meetings will be accessible to persons with disabilities. An assistive listening system, computer assisted real-time transcription (CART), and sign language interpreters will be provided. Persons attending the meetings are requested to refrain from using perfume, cologne, and other fragrances for the comfort of other participants (*see* www.access-board.gov/about/policies/fragrance.htm for more information). Also, persons wishing to provide handouts or other written information to the committee are requested to provide electronic formats to Rex Pace via email prior to the meetings so that alternate formats can be distributed to committee members.

David M. Capozzi,
Executive Director.

[FR Doc. 2013-00071 Filed 1-7-13; 8:45 am]

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FEDERAL COMMUNICATIONS COMMISSION

47 CFR Parts 1 and 27

[WT Docket No. 12-357; FCC 12-152]

Service Rules for the Advanced Wireless Services in the H Block— Implementing Section 6401 of the Middle Class Tax Relief and Job Creation Act of 2012 Related to the 1915-1920 MHz and 1995-2000 MHz Bands

AGENCY: Federal Communications Commission.

ACTION: Notice of proposed rulemaking.

SUMMARY: In this document, the Commission proposes rules for the Advanced Wireless Services (AWS) H Block that would make available ten megahertz of spectrum for flexible use. The proposal would extend the widely-deployed Personal Communications Services (PCS) band, which is used by the four national providers as well as regional and rural providers to offer mobile service across the nation. The additional spectrum for mobile use will help ensure that the speed, capacity, and ubiquity of the nation's wireless networks keeps pace with the skyrocketing demand for mobile service.

DATES: Submit comments on or before February 6, 2013. Submit reply